



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶: A61L 15/46, 15/60	A1	(11) International Publication Number: WO 98/57677 (43) International Publication Date: 23 December 1998 (23.12.98)
(21) International Application Number: PCT/SE98/01111 (22) International Filing Date: 10 June 1998 (10.06.98) (30) Priority Data: 9702298-2 17 June 1997 (17.06.97) SE (71) Applicant (for all designated States except US): SCA MÖLNLYCKE AB [SE/SE]; S-405 03 Göteborg (SE). (72) Inventors; and (75) Inventors/Applicants (for US only): RUNEMAN, Bo [SE/SE]; Jons Väg 6, S-433 75 Jonsered (SE). FORSGREN-BRUSK, Ulla [SE/SE]; Plommonvägen 35, S-435 43 Pixbo (SE). (74) Agents: GRAUDUMS, Valdis et al.; Albihns Patentbyrå Göteborg AB, P.O. Box 142, S-401 22 Göteborg (SE).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>In English translation (filed in Swedish).</i>
(54) Title: REDUCTION OF UNWANTED SIDE-EFFECTS DURING USE OF ABSORBENT ARTICLES BY MEANS OF pH-CONTROL (57) Abstract <p>The present invention relates to absorbent articles such as diapers, incontinence protectors, sanitary napkins, wound dressings and similar articles which are applied in contact with skin, and concerns methods which make it possible to use an article for an extended period of time without the appearance of unwanted side-effects such as, for instance, growth of undesirable micro-organisms. The absorbent body in the absorbent article comprises a pH-controlling substance in the form of a partially neutralised super-absorbent material in such a manner that the pH in the absorbent article after wetting is in the interval 3.5 - 4.9. Growth of undesirable strains of micro-organisms is prevented and unwanted side-effects resulting from use of the article are reduced.</p>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	R	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

TITLE:

5

REDUCTION OF UNWANTED SIDE-EFFECTS DURING USE OF ABSORBENT
ARTICLES BY MEANS OF pH-CONTROL

TECHNICAL FIELD:

10

The present invention relates to absorbent articles such as
diapers, incontinence protectors, sanitary napkins, wound
dressings and similar articles which are applied against
skin, and concerns methods aiming at the reduction of
15 unwanted side-effects which sometimes occur during use of
said articles.

BACKGROUND:

20

Absorbent articles of this kind are known in a number of
different types. Conventionally, the absorbent body in
these articles is produced by dry-defibration of cellulose
pulp from, for instance, rolls, bales, or sheets and
converting the pulp into a web of fluffed pulp sometimes
25 with the admixture of what is known as superabsorbents,
which are polymers with the ability to absorb several times
their own weight of water or body fluids.

30

All uses of products which are applied against skin may
lead to unwanted side-effects. These may occur as a result
of occlusion, moisture, mechanical, microbial, and
enzymatic factors which all, to different degrees, interact
and amplify the influence of each other and may cause
different forms of skin irritation and primary or secondary
35 skin infections which sometimes occur in users of said
articles. An increase in pH is a normal phenomenon during
use of absorbent articles in contact with skin. However,
several unwanted side-effects may occur as a result of, or
in connection with, a pH-increase. One example of an
40 unwanted side-effect of this kind is irritational contact

dermatitis which exhibits a connection with the surface-pH of skin. Factors Predisposing Cutaneous Irritation, Wilhelm K-P et al, Dermatologic Clinics 8, No. 1, 1990.

5 Another example of unwanted effects is the activity of enzymes such as lipases and proteases which exhibit a strongly pH-dependent activity which increases with increasing pH. The skin starts to decompose and becomes sensitive to mechanical forces and bacterial attacks.
10 Etiologic Factors in Diaper Dermatitis: The Role of Urine, Berg R.W. et al, Pediatric. Dermatology. 3, No. 2, 1986.

Another example of unwanted side-effects is that some bacteria such as Proteus can metabolise substances in urine
15 and other body fluids and produce odorous substances such as ammonia and amines, which also raises the pH. At a high pH, the equilibrium for many odorous substances is shifted so that more volatile components are formed, resulting in a stronger odour than at a low pH.

20 An absorbent article such as a diaper, or the like provides an environment for micro-organisms which comprises access to moisture, nutrients, and heat. An unwanted side-effect of this is that growth of bacteria is promoted in such an
25 environment. High numbers of bacteria constitute a risk of the appearance of infections. Moreover, a high presence of bacteria constitutes an increased risk for the creation of unpleasant odours caused by the formation of different substances, resulting from biological or chemical
30 decomposition of components in body fluids such as, for instance, urine or menses.

Micro-organisms which may be associated with the occurrence of problems when using products in contact with skin may be
35 of different types. Examples of micro-organisms which cause odour and those which constitute a risk for urinary tract

infections are *Proteus mirabilis*, *Proteus vulgaris*, *Echerichia coli*, *Enterococcus* and *Klebsiella*.

5 Examples of micro-organisms which are associated with skin-infections and other skin problems are *Candida albicans* and *Staphylococcus* sp. and *Streptococcus* sp.

RELATED ART:

10 It is known that a low pH is advantageous in order to reduce the occurrence of negative effects on skin.

15 Different ways of solving the described problems have been suggested. In the American patent US 3,794,034 the importance of the pH in an absorbent article is described as well as the impregnation of an absorbent article with buffering substances which aid in keeping the pH in the article between 3.5 and 6.0, which is advantageous both for inhibiting the growth of unwanted bacteria and, in
20 connection therewith, the creation of unwanted odours, and for avoiding negative effects on skin.

25 In the European patent 0,311,344 pH-control in an absorbent article is disclosed, wherein the buffering properties are obtained by using a partially neutralised superabsorbent material. In order to achieve sufficient inhibition of the growth of unwanted bacteria in the article, a separate bacteria inhibiting substance has also been added. The European patent EP 0,316,518 discloses buffering of pH in
30 an absorbent article by using a polymeric organic acid.

35 One drawback in using bacteria suppressing agents, as described in EP 0,311,344, is that these agents are often selective and may be associated with risks, for instance in the form of allergenic properties or negative ecological consequences in garbage handling. Furthermore, the use of

this type of agents may involve a risk that resistant strains of micro-organisms arise.

5 In EP 0,316,518 an absorbent article is disclosed, comprising a pH-controlling substance in the form of a partially neutralised superabsorbent material. It is stated that pH in the absorbent article is in the interval 5 - 6. However, this is not a sufficiently low pH in order to
10 obtain a sufficient inhibition effect on the micro-organisms which are present since the majority of micro-organisms require a pH below 5 in order to be affected to a significant degree.

15 One difficulty when evaluating the influence of different components on the pH in absorbent articles of the aforementioned kind is that the degree of acidity in cellulose fluff pulp varies depending on the production method. Chemical cellulose pulp (CP) varies between pH 6 - 8.5 and chemical thermomechanical cellulose pulp (CTMP)
20 varies between 5.5 - 8.5. Variations outside said intervals also occur.

OBJECT OF THE INVENTION:

25 The object of the present invention is to achieve an absorbent article of the kind mentioned in the introduction which makes it possible to use the article during a longer period of time without the appearance of undesired side effects such as unpleasant odour, increased risk of
30 infections or negative skin effects which are due to the growth of micro-organisms resulting from an unwanted raise in skin-pH, or from other reasons.

DESCRIPTION OF THE INVENTION:

An absorbent article of the kind mentioned in the introduction, wherein the problems connected with using the article for an extended period of time have been substantially removed, has been achieved through the invention with an absorbent body in the absorbent article comprising a pH-controlling substance in the form of a partially neutralised superabsorbent material and in that pH in the absorbent article after wetting is in the interval 3.5 - 4.9, whereby growth of unwanted strains of micro-organisms is restricted and the occurrence of unwanted side effects resulting from the use of the article is reduced.

It has been shown that if the absorbent body in an absorbent article comprises a pH-controlling substance in the form of a partially neutralised superabsorbent material which after wetting creates a pH in the absorbent article which is in the interval 3.5 - 4.9, and preferably 4.1 - 4.7, a significant inhibition effect on the growth of unwanted micro-organisms is obtained. The inhibition effect is based on the fact that many micro-organisms have an activity which is strongly pH-dependent and decreases with decreasing pH, which means that a decrease in pH leads to a decrease in activity in most micro-organisms which, in turn, leads to a decrease of bad smell as well as negative effects on skin in the form of skin-irritation and primary or secondary skin-infections and a generally lower risk of infections.

Enzymes, such as lipases and proteases have an activity which is strongly pH-dependent and decreases with decreasing pH, which means that a decrease in pH will also lead to a decrease in enzymatic activity and an accompanying reduction in negative skin effects.

An absorbent body containing absorbent material and absorbed fluid is a heterogeneous system from a pH point of view. The system may contain superabsorbent material, fibres and liquid containing several kinds of ions. In order to obtain reproducible pH-values, measurements must be made in several places in the absorbent body and the mean value calculated.

An absorbent body in accordance with the invention may also comprise other absorbent materials besides a partially neutralised superabsorbent material, for instance fluffed cellulose pulp. It has proven advantageous to use a partially neutralised superabsorbent material as above in combination with fluffed cellulose pulp having a pH below 7, preferably below 6 which gives a further improved effect.

A suitable fluffed cellulose pulp may consist of a chemical thermo-mechanical cellulose pulp having a pH = 2.5 - 8.5, preferably 2.5 - 6.5 and most preferably 2.5 - 5.5, or of a chemical cellulose pulp having a pH = 2.5 - 8.5, preferably 2.5 - 8.0 and most preferably 2.5 - 7.0.

In order to obtain a suitable degree of acidity in the cellulose pulp, its pH may be controlled during the production process, for instance by adding an acidifying substance. This added substance may, for instance, consist of SO₂-water. In this way, it is also possible to neutralise NaOH which may occur in the pulp. Another way of achieving a suitable degree of acidity in the cellulose pulp is to add a suitable acid after production of the pulp.

A suitable, partially neutralised superabsorbent material may, for instance, consist of a cross-linked poly-acrylate of the kind which is described in the European patent EP 0

391 108 in the name of Cassella AG. Other types of
superabsorbent material than the above indicated, and
having corresponding properties, may be used. A suitable
proportion of superabsorbent material in the article is 5 -
5 100%, preferably 15-50%, and most preferably 15-50%. The
proportion of superabsorbent material which has been stated
to be preferred relates to areas of the article which will
absorb the major part of the fluid and, consequently, does
not concern end-portions or edge-portions or similar parts
10 of the article since such parts thereof mostly do not
significantly contribute to the absorptive function of the
article.

Examples of the relation between the degree of
15 neutralisation and pH in the superabsorbent material are
given below.

	Degree of neutralisation %	pH
20	18	4.0
	25	4.3
	30	4.5
	35	4.7
	45	5.0
25	60	5.5

From the table it can be learned that the degree of
neutralisation should normally be lower than 45% and
preferably 35%. However, the degree of neutralisation
30 should preferably be higher than approximately 20%.

An absorbent body in accordance with the invention,
comprising a partially neutralised super-absorbent material
in accordance with the invention may have somewhat lower
35 absorption capacity when compared to a corresponding
absorbent body containing a conventional super-absorbent

material. Such a lowering of the absorption capacity may be compensated by increasing the amount of absorbent material to a corresponding degree.

5 An absorbent body in accordance with the invention, comprising a partially neutralised super-absorbent material may also comprise some type of conventional bacteria-inhibiting substance such as parabens or benzoic acid. Such bacteria inhibitors normally exhibit an increased effect at
10 a lower pH.

SHORT DESCRIPTION OF DRAWINGS:

The invention will in the following be described in more
15 detail, with reference to a number of examples which are shown in the appended figures.

Fig. 1 shows in a diagram the formation of ammonia in a reference product compared with a product in accordance with the invention.
20

Fig. 2 shows in a diagram the surface-pH of the skin during use of a test product containing a conventional absorption body, compared with the use of a corresponding test product in accordance with the invention.
25

30 DESCRIPTION OF EXAMPLES:

The following examples are intended to give a closer illustration of the effect in absorbent articles having an absorbent body comprising a combination of a partially
35 neutralised super-absorbent material and cellulose pulp

having a pH of 2.5 - 8.5 compared to conventional materials of a corresponding type.

TEST LIQUIDS:

5

Test liquid 1

A solution of 0.9% sodium chloride.

10

Test liquid 2

Synthetic urine according to the description in, among other, EP 0,565,606 which can be obtained from Jayco Pharmaceuticals Co., Pennsylvania. The composition is 2 g/l KCl; 2 g/l Na₂SO₄; 0.85 g/l (NH₄)H₂PO₄; 0.15 g/l (NH₄)₂HPO₄; 0.19 g/l CaCl₂ and 0.23 g/l MgCl₂. The pH in this composition is 6.0 - 6.4.

20

Test liquid 3

Synthetic urine containing the following substances: KCl, NaCl, MgSO₄, KH₂PO₄, Na₂HPO₄, NH₂CONH₂. The pH in this composition is 6.0 - 6.5.

25

Test liquid 4

Sterile synthetic urine to which has been added a growth medium for micro-organisms. The synthetic urine contains mono- and divalent cat- and anions and urea and has been prepared in accordance with the information in Geigy, Scientific Tables, vol 2, 8th ed. 1981 p. 53. The growth medium for micro-organisms is based on information of Hook- and FSA-media for entero-bacteria. The pH in this mixture is 6.6.

35

TEST METHODS:

Method 1, preparation of absorbent bodies for test.

5 Absorbent bodies were prepared using a slightly modified
sample former according to SCAN C 33:80. Fluffed pulp and
super-absorbent material of the desired type were weighed
and a homogeneous mixture of fluffed pulp and super-
10 absorbent material was subsequently introduced into a flow
of air having a negative pressure of approximately 85 mbar,
through a pipe having a diameter of 5 cm and being equipped
at the bottom with a metal net having a thin tissue placed
thereon. The mixture of fluffed pulp and super-absorbent
15 material was gathered onto the tissue on the metal net and
thereafter constituted the absorbent body. The absorbent
body was weighed and compressed to a bulk of 6-12 cm³/g. A
number of absorbent bodies referred to as Reference product
1, Reference product 2, test product 1, test product 2,
20 test product 3, test product 4, etc. having different
compositions as described below were made. The amount of
absorbent material in the single core and dual core
absorbent bodies, respectively, was adjusted so that the
single cores and dual cores had approximately the same
absorption capacity.

25

Method 2, measurement of pH in cellulose pulp.

The pH in the cellulose pulp in the different test products
was measured by determining the pH in a water extract from
30 the pulp in accordance with SCAN P 14:65. 1.0 g air dry
cellulose pulp was placed in a 100 ml glass container and
20 ml distilled water was added. After mixing, a further 50
ml of distilled water was added and the mixture was stirred
for approximately 30 s and was left for 1 hour. The liquid
35 was poured off and pH was determined with a glass electrode

at 20-30°C. Two samples were prepared and the mean value was calculated.

Method 3, measurement of pH in an absorbent body.

5

An absorbent body having a diameter of approximately 50 mm was prepared according to method 1. A predetermined amount of Test liquid 1, 2 or 3 was added, 10 ml to a single core absorbent body and 20 ml to a dual core absorbent body, 10 whereafter the absorbent body was left to swell for 30 minutes. Thereafter, pH was measured in the absorbent body using a surface electrode, Flat-bottomed Metrohm pH-meter, Beckman Ø12 or Ø72. Parallel measurements were performed on at least two different absorbent bodies. The pH was 15 determined at 10 locations on each absorbent body and the mean value was calculated.

Method 4, measurement of bacteria inhibition in absorbent bodies.

20

Absorbent bodies were prepared in accordance with method 1. Single core, as well as dual core absorbent bodies were prepared. Test liquid 4 was prepared. Bacteria suspensions of each of Escherichia coli (E.c.), Proteus mirabilis 25 (P.m.), Enterococcus faecalis (E.f.) were cultivated in nutritional bouillon 30°C overnight. The graft cultures were diluted and the bacterial count was determined. The cultures were mixed in different proportions so that the final mixed culture contained approximately 10⁴ organisms 30 per ml test liquid 4. Test liquid 4 was added to a sterile sputum container 70.5 x 52 mm, volume 100 ml, and the absorbent body was placed upside-down in the container and was left to absorb liquid for 5 minutes, whereafter the container was turned and incubated at 35 °C for 0; 6 and 12 35 hours, respectively whereafter the bacterial count in the absorbent body was determined. The nutritional medium used

12

was TGE agar for measurement of the total amount of bacteria and Drigalski agar for specific determination of *Escherichia coli* and *Proteus mirabilis*, and Slanetz Bartley agar for specific determination of *Enterococcus faecalis*.

5

Method 5, measurement of ammonia content.

Single core absorbent bodies were prepared in accordance with method 1. Test liquid and micro-organisms were added in accordance with method 5 whereafter the containers were incubated at 35°C for 0; 3; 6 and 12 hours, whereafter samples were taken from the containers using a hand pump and a so called Dräger-pipe. The ammonia content was obtained as a colour change on a scale graded in ppm or volume-percent.

15

Method 6, measurement of the surface-pH of skin.

Test products were prepared by applying a backing of approximately 25 g/m² polyethylene and a topsheet of approximately 20 g/m² polypropylene nonwoven to absorbent bodies according to Ref. 3 and test 4, respectively. Test liquid 3 was added to the topsheet and was absorbed into the test product. The test products which were obtained in this manner were applied to the forearms of a test person and were left there for 24 hours. The procedure was repeated twice. The surface-pH of the skin at the place of contact was measured before application and after 24, 48 and 72 hours with Courage + Khazaka skin-pH-meter with a flat-bottomed Mettler-Toledo glass electrode 403/120.

25

30

TEST PRODUCTS:

Reference product 1: Single core absorbent body having a total weight of 1 gram, prepared from a conventional super-absorbent

35

material and a conventional chemical thermo-mechanical cellulose pulp with the ratio 15/85 weight-%.

5 Test product 1: Single core absorbent body having a
total weight of 1 gram, prepared from
a partially neutralised super-
absorbent material with pH = 4.2, in
accordance with the invention, and a
10 chemical thermo-mechanical cellulose
pulp with pH = 5.8 and with the ratio
15/85 weight-%.

15 Test product 2: Single core absorbent body having a
total weight of 1 gram, prepared from
a partially neutralised super-
absorbent material with pH = 4.2, in
accordance with the invention, and a
20 chemical thermo-mechanical cellulose
pulp with pH = 3.7 and with the ratio
15/85 weight-%.

Reference product 2: Dual core absorbent body. The upper core (uc) had a total weight of 1.2 grams and was prepared from a conventional super-absorbent material and a conventional chemical thermo-mechanical cellulose pulp with the ratio 12/88 weight-%. The lower core (lc) had a total weight of 1.1 grams and was prepared from a conventional super-absorbent material and a conventional chemical cellulose pulp with the ratio 12/88 weight-%.

Test product 3: Dual core absorbent body. The upper core (uc) had a total weight of 1.3 grams and was prepared from a partially neutralised super-absorbent material having a pH = 4.5, in accordance with the invention, and a chemical thermo-mechanical cellulose pulp having a pH = 5.8 and with the ratio 15/85 weight-%. The lower core (lc) had a total weight of 1.2 grams and was prepared from a partially neutralised super-absorbent material having a pH = 4.5, in accordance with the invention, and a chemical cellulose pulp having a pH = 6.3 and with the ratio 15/85 weight-%.

Reference product 3: Single core absorbent body having a total weight of 1 gram, prepared from a conventional super-absorbent material and a conventional chemical cellulose pulp, with the ratio 15/85 weight-%.

Test product 4: Single core absorbent body having a total weight of 1 gram, prepared from a partially neutralised super-absorbent material with pH = 4.2, in accordance with the invention, and a conventional chemical cellulose, with the ratio 15/85 weight-%.

Reference product 4: Single core absorbent body having a total weight of 1 gram, prepared from a conventional super-absorbent material and a chemical thermo-

15

mechanical cellulose pulp with pH = 6.7, with the ratio 15/85 weight-%.

Test product 5:

5

Single core absorbent body having a total weight of 1 gram, prepared from a partially neutralised super-absorbent material with pH = 4.2, in accordance with the invention, and a chemical thermo-mechanical cellulose pulp with pH = 6.7 and with the ratio 15/85 weight-%.

10

Test product 6:

15

Dual core absorbent body. The upper core (uc) had a total weight of 1.3 grams and was prepared from a partially neutralised super-absorbent material having a pH = 4.6, in accordance with the invention, and a chemical thermo-mechanical cellulose pulp having a pH = 5.8 and with the ratio 15/85 weight-%. The lower core (lc) had a total weight of 1.2 grams and was prepared from a partially neutralised super-absorbent material having a pH = 4.6, in accordance with the invention, and a chemical cellulose pulp having a pH = 6.3 and with the ratio 15/85 weight-%.

20

25

30 TEST RESULTS:

Example 1

35

Table 1 shows that in a single core, conventional absorbent body according to reference product 1, good growth of

micro-organisms prevails. The measurement was performed in accordance with Method 4.

Table 1:

time	Esherichia coli	Proteus mirábilis	Enterococcus faecalis
0 hours	3,3	3,1	3,7
6 hours	7,0	6,4	7,1
12 hours	9,2	9,1	8,3

10 Example 2

Table 2 shows that in a single core absorbent body according to test product 1, good inhibition of the growth of micro-organisms is obtained. The measurement was performed in accordance with Method 4.

Table 2:

time	Esherichia coli	Proteus mirábilis	Enterococcus faecalis
0 hours	3,2	3,3	3,4
6 hours	5,5	3,2	4,8
12 hours	7,3	4,0	6,1

20 Example 3

Table 3 shows that in a single core absorbent body according to test product 2, good inhibition of the growth of micro-organisms is obtained. The measurement was performed in accordance with Method 4.

Table 3:

time	Esherichia coli	Proteus mirabilis	Enterococcus faecalis
0 hours	3,4	3,3	3,5
6 hours	3,2	2,6	3,6
12 hours	2,8	2,0	3,5

Example 4

Table 4 shows that in a dual core, conventional absorbent body according to reference product 2, good growth of micro-organisms prevails. The measurement was performed in accordance with Method 4.

Table 4:

time	Esherichia coli		Proteus mirabilis		Enterococcus faecalis	
	uc*	lc**	uc*	lc**	uc*	lc**
0 hrs	3,4	3,4	3,4	3,4	3,4	3,4
6 hrs	6,8	7,0	6,6	6,7	6,7	6,2
12 hrs	9,0	9,0	9,1	9,0	8,0	7,8

*uc = upper core, **uk = lower core

Example 5

Table 5 shows that in a single core, absorbent body according to test product 3, good inhibition of the growth of micro-organisms is obtained. The measurement was performed in accordance with Method 4.

Table 5:

time	Esherichia coli		Proteus mirábilis		Enterococcus faecalis	
	uc*	lc**	uc*	lc**	uc*	lc**
0 hrs	3,4	3,4	3,4	3,4	3,4	3,4
6 hrs	5,1	5,6	3,3	4,2	4,4	4,5
12 hrs	7,3	7,4	4,0	4,0	5,9	4,8

*uc = upper core, **lc = lower core

Example 6

Fig. 1 shows that efficient delay of the development of ammonia is obtained in a single core absorbent body according to test product 5 when compared to a single core, conventional absorbent body, according to Reference product 4. The measurement was performed in accordance with Method 5.

Example 7

Fig. 2 shows that the surface-pH of skin after a period of use of a test product containing an absorbent body in accordance with the invention, test product 4, is stabilised at a lower level than after use of a corresponding test product containing a conventional super-absorbent material, according to Reference product 3, after addition of Test liquid 3. The measurement was performed in accordance with Method 6.

Example 8

Table 6 shows that pH, when measured in a single core absorbent body, test product 1, in accordance with the invention, after addition of test liquid, lies within the effective pH-interval 3.5 - 4.9. The measurement was performed in accordance with Method 3.

Table 6:

	Test liquid 1	Test liquid 2	Test liquid 3
pH	4,29	4,42	4,54

Example 9

Table 7 shows that pH, when measured in a dual core absorbent body, test product 6, in accordance with the invention, after addition of test liquid, lies within the effective pH-interval 3.5 - 4.9. The measurement was performed in accordance with Method 3.

Table 7:

	Test liquid 1	Test liquid 2	Test liquid 3
pH uc*	4,72	4,83	4,80
pH lc*	4,75	4,73	4,73

*uc = upper core, **lc = lower core

The invention shall not be considered to be restricted to the embodiments described herein. Accordingly, a number of further variants and modifications are conceivable within the scope of the appended claims.

CLAIMS:

5

1. An absorbent article, intended to be worn in contact with the skin of a wearer, said article comprising a pH-controlling substance in the form of a partially neutralised super-absorbent material,

10

characterized in that the pH in the article, after wetting and during use of the article in contact with skin, is in the interval 3.5 - 4.9, preferably 4.1 - 4.7.

15

2. An absorbent article according to claim 1, characterized in that said article comprises at least one further absorbent material.

20

3. An absorbent article according to claim 2, characterized in that the article comprises chemical thermo-mechanical cellulose pulp (CTMP).

25

4. An absorbent article according to claim 3, characterized in that the chemical thermo-mechanical cellulose pulp has a pH from 2.5 - 8.5, preferably 2.5 - 6.5 and most preferably 2.5 - 5.5.

30

5. An absorbent article according to claim 4, characterized in that said pH has been obtained by the addition of an acidifying agent.

35

6. An absorbent article according to claim 4, characterized in that said pH has been obtained by the addition of an acidifying agent in a separate step after the pulp manufacturing process.

40

7. An absorbent article according to any one of claims 2-6, characterized in that the article comprises chemical cellulose pulp (CP).

8. An absorbent article according to claim 7,

21

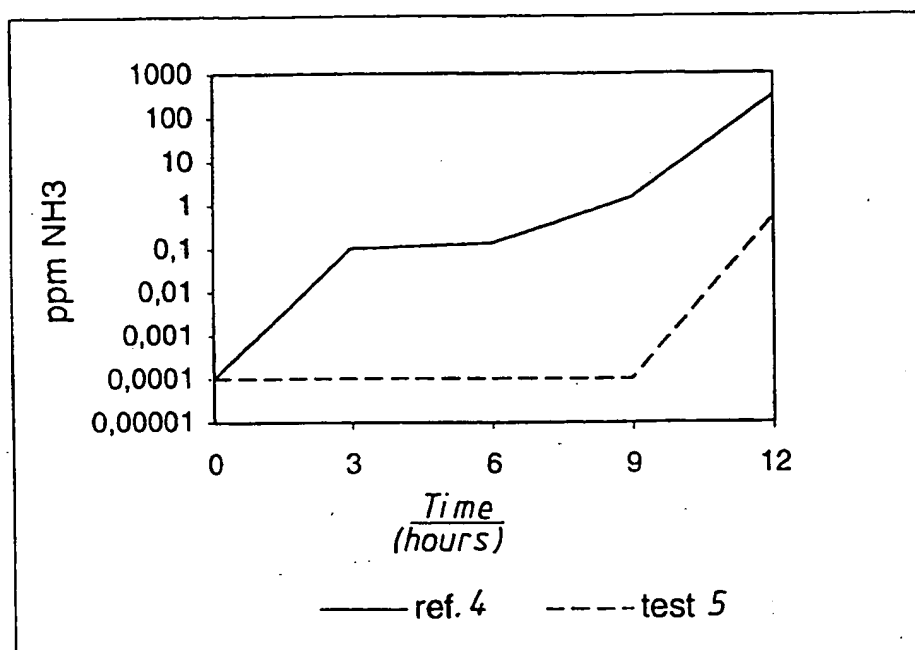
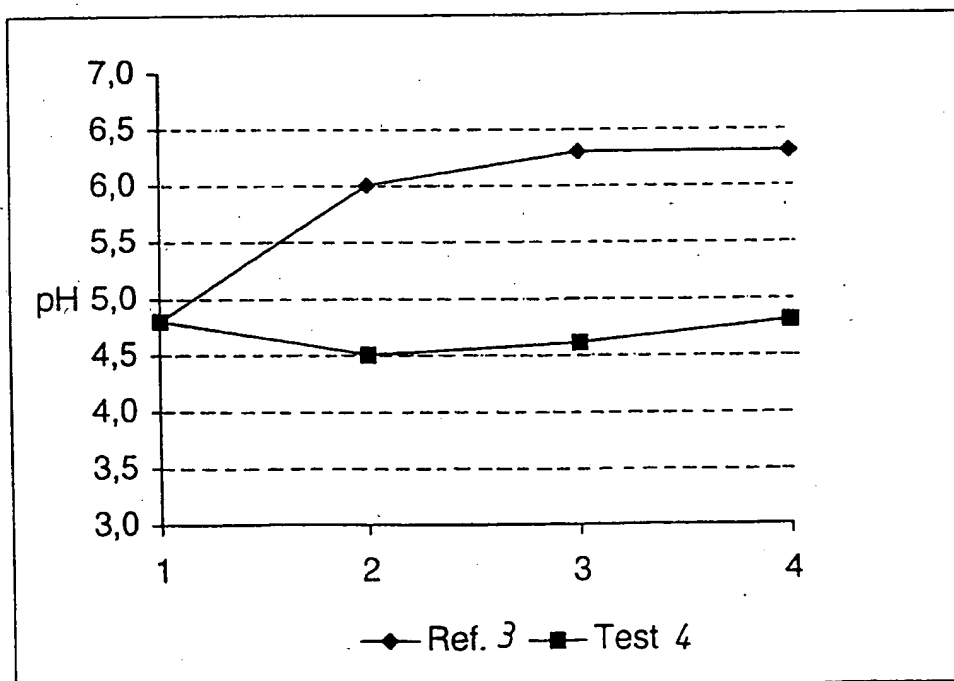
c h a r a c t e r i z e d i n that said chemical cellulose pulp has a pH fr m 2.5 - 8.5, preferably 2.5 - 8.0 and most preferably 2.5 - 7.0.

5 9. An absorbent article according to claim 1,
c h a r a c t e r i z e d i n that said pH has been obtained in the pulp manufacturing process by the addition of an acidifying agent.

10 10. An absorbent article according to claim 1,
c h a r a c t e r i z e d i n that said pH has been obtained by the addition of an acidifying agent in a separate step after the pulp manufacturing process.

15

1/1

FIG. 1FIG. 2

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 98/01111

A. CLASSIFICATION OF SUBJECT MATTER

IPC6: A61L 15/46, A61L 15/60

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: A61L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0202126 A2 (THE PROCTER & GAMBLE COMPANY), 20 November 1986 (20.11.86), page 7, line 1- line 13; example 9	1-2
Y	--	3-10
Y	WO 9315702 A1 (MÖLNLYCKE AB), 19 August 1993 (19.08.93), page 7, line 28 - page 8, line 23; abstract	3,7
Y	US 3794034 A (JOHN LESLIE JONES, SR.), 26 February 1974 (26.02.74), column 3, line 66 - column 4, line 15; claim 1	4-6,8-10
	-- -----	

☐ Further documents are listed in the continuation of Box C.☒ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

8 Sept 1998

Date of mailing of the international search report

21 -09- 1998

Name and mailing address of the ISA/
Swedish Patent Office
Box 5055, S-102 42 STOCKHOLM
Facsimile No. +46 8 666 02 86

Authorized officer

Jack Hedlund
Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT
Information on patent family members

27/07/98

International application No.

PCT/SE 98/01111

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0202126 A2	20/11/86	SE 0202126 T3	
		AT 113482 T	15/11/94
		AU 577016 B	08/09/88
		AU 5741686 A	20/11/86
		CA 1259175 A	12/09/89
		DE 3650117 D,T	24/05/95
		DK 169694 B	16/01/95
		DK 226386 A	16/11/86
		EG 17694 A	30/10/90
		FI 87310 B	15/09/92
		FI 862009 A	16/11/86
		GB 2175211 A,B	26/11/86
		HK 10692 A	31/01/92
		IE 64373 B	26/07/95
		JP 62033804 A	13/02/87
		KR 9401377 B	21/02/94
		MX 168802 B	09/06/93
		PT 82572 B	03/03/88
		US 4657537 A	14/04/87
WO 9315702 A1	19/08/93	CA 2117342 A	19/08/93
		DE 69121399 D,T	12/12/96
		DK 625895 T	20/04/98
		EP 0525090 A,B	03/02/93
		SE 0525090 T3	
		EP 0625895 A,B	30/11/94
		SE 0625895 T3	
		SE 468744 B,C	15/03/93
		SE 9100274 A	30/07/92
		SK 402292 A	09/08/95
		US 5343898 A	06/09/94
US 3794034 A	26/02/74	NONE	